Application No.: 10/568,992 Atty. Docket No.: 032301.443

Response dated August 2, 2011

Reply to Final Office Action of May 10, 2011

AMENDMENT TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

Please amend the claims as follows:

 $1. \ (Currently \ amended) \ Surface-modified \ pyrogenically \ produced, \ \underline{nano-scale} \ zinc \ oxide \ powder, \ comprising \ aggregates \ and \ having the following \ physico-chemical \ characteristic \ data:$

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BET surface areas: 18 ± 5 m²/g
C content: 0.5 to 1.0 wt.%,
average diameter: 50 to 300 nm,
shape factor F (circle) of below 0.5,
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 $wherein \ the \ surface \ modification \ includes \ silanization \ with \ a \ surface \ modifying \ agent \ selected \ from \ the \ group \ consisting \ of:$

- a) Organosilanes of the type (RO)₃Si(C_nH_{2n+1}) and RO)₃Si(C_nH_{2n-1}) R = alkyl, such as, methyl-, ethyl-, n-propyl-, i-propyl-, butyl- n = 1 -20
- b) Organosilanes of the type $R'_x(RO)_ySi(C_nH_{2n+1})$ and $R'_x(RO)_ySi(C_nH_{2n-1})$ R= alkyl, such as, methyl-, ethyl-, n-propyl-, i-propyl-, butyl- R'= alkyl, such as, methyl-, ethyl-, n-propyl-, i-propyl-, butyl- R= cycloalkyl n=1-20 x+y=3 x=1,2 y=1,2
- c) Halogeno-organosilanes of the type $X_3Si(C_nH_{2n+1})$ and $X_3Si(C_nH_{2n-1})$ X = Cl, Br $_n$ = 1 20
- d) Halogeno-organosilanes of the type $X_2(R')Si(C_nH_{2n+1})$ and $X_2(R')Si(C_nH_{2n-1})$ X = Cl, Br R' = alkyl, such as, methyl-, ethyl-, n-propyl-, i-propyl-, butyl-

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R'=cycloalkyl

n = 1 - 20
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e) Halogeno-organosilanes of the type

$$\begin{split} &X(R')_2Si(C_nH_{2n+1}) \text{ and } X(R')_2Si(C_nH_{2n-1})\\ &X=Cl, Br\\ &R'=\text{alkyl, such as, methyl-, ethyl-, n-propyl-, i-propyl-, butyl-}\\ &R'=\text{cycloalkyl}\\ &n=1-20 \end{split}$$

f) Organosilanes of the type (RO)3Si(CH2)m-R'

R = alkyl, such as methyl-, ethyl-, propyl-

m = 0,1 - 20

R' = methyl-, aryl (-C₆H₅, substituted phenyl radicals)

-N-(CH2-CH2-NH2)2

 $-OOC(CH_3)C = CH_2$

-OCH2-CH(O)CH2

-NH-CO-N-CO-(CH2)5

-NH-COO-CH3, -NH-COO-CH2-CH3, -NH-(CH2)3Si(OR)3

 $-\mathrm{S}_{x}\text{-}(\mathrm{CH}_{2})_{3}\mathrm{Si}(\mathrm{OR})_{3}$

-SH -NR'R"R"" (R' = alkyl, aryl; R" = H,

alkyl, aryl; R''' = H, alkyl, aryl, benzyl, C₂H₄NR'''' R''''' where R'''' = H, alkyl and

 $R^{""} = H$, alkyl)

g) Organosilanes of the type (R")x(RO)vSi(CH2)m-R'

$$R'' = alkyl \quad x+y = 2$$

= cycloalkyl x = 1, 2

y = 1, 2

m = 0,1 to 20

R' = methyl-, aryl (-C₆H₅, substituted phenyl radicals)

-C₄F₉, -OCF₂-CHF-CF₃, -C₆F₁₃, -O-CF₂-CHF₂ -NH₂, -N₃, -SCN, -CH=CH₂, -NH-CH₂-CH₂-NH₂,

-N-(CH2-CH2-NH2)2

 $-OOC(CH_3)C = CH_2$

-OCH₂-CH(O)CH₂

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-NH-CO-N-CO-(CH2)5
             -NH-COO-CH3, -NH-COO-CH2-CH3, -NH-(CH2)3Si(OR)3
             -Sx-(CH2)3Si(OR)3
             -SH
             - NR'R"R" (R' = alkyl, aryl; R" = H, alkyl, aryl; R" = H, alkyl, aryl,
                  C_2H_4NR''''R''''' where R''''=H, alkyl
                  and R"" = H, alkyl)
h) Halogeno-organosilanes of the type X3Si(CH2)m- R'
        X = Cl. Br
        m = 0.1 - 20
        R' = methyl-, aryl (-C<sub>6</sub>H<sub>5</sub>, substituted
           phenyl radicals)
           -C4F9, -OCF2-CHF-CF3, -C6F13, -O-CF2-CHF2
           -NH2, -N3, -SCN, -CH=CH2,
           -NH-CH2-CH2-NH2
           -N-(CH2-CH2-NH2)2
           -OOC(CH_3)C = CH_2
           -OCH2-CH(O)CH2
           -NH-CO-N-CO-(CH2)5
           -NH-COO-CH3, -NH-COO-CH2-CH3, -NH-(CH2)3Si(OR)3
           -Sx-(CH2)3Si(OR)3
          -SH

 Halogeno-organosilanes of the type (R)X2Si(CH2)m-R<sup>t</sup>

        X = Cl. Br
        R = alkyl, such as methyl,- ethyl-, propyl-
        m = 0.1 - 20
        R' = methyl-, aryl (-C<sub>6</sub>H<sub>5</sub>, substituted phenyl radicals)
           -C4F9, -OCF2-CHF-CF3, -C6F13, -O-CF2-CHF2
           -NH2, -N3, -SCN, -CH=CH2, -NH-CH2-CH2-NH2,
           -N-(CH2-CH2-NH2)2
           -OOC(CH_3)C = CH_2
           -OCH2-CH(O)CH2
           -NH-CO-N-CO-(CH2)5
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-NH-COO-CH₃, -NH-COO-CH₂-CH₃, -NH-(CH₂)₃Si(OR)₃, wherein R can be methyl-, ethyl-, propyl-, butyl-

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 $-S_X\text{-}(CH_2)_3Si(OR)_3,$ wherein R can be methyl-, ethyl-, propyl-, butyl-SH

j) Halogeno-organosilanes of the type (R)2X Si(CH2)m-R'

k) Silazanes of the type $R'R_2Si-N-SiR_2R'$

R' = alkyl, vinyl, aryl

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 Cyclic polysiloxanes of the type D 3, D 4, D 5, wherein D 3, D 4 and D 5 are understood as cyclic polysiloxanes with 3, 4 or 5 units of the type -O-Si(CH₃)₂-.E.g. octamethylcyclotetrasiloxane = D 4

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m) Polysiloxanes or silicone oils of the type

$$\begin{array}{c} \text{Y-O-} & \left(\begin{matrix} R \\ | \\ | \\ | \\ \text{Si-O} \\ | \\ R' \end{matrix} \right)_{m} - \left(\begin{matrix} R'' \\ | \\ | \\ \text{Si-O} \\ | \\ R' \end{matrix} \right)_{n} - \left(\begin{matrix} R'' \\ | \\ | \\ \text{Si-O} \\ | \\ \text{-Y} \end{matrix} \right)_{u} - \left(\begin{matrix} m = 0,1,2,3,...\infty \\ n = 0,1,2,3,...\infty \\ n = 0,1,2,3,...\infty \\ u = 0,1,2,3,...\infty \\ u = 0,1,2,3,...\infty \\ u = 0,1,2,3,...\infty \\ u = 0,1,2,3,...\infty \\ \text{New } \left(\begin{matrix} n = 0,1,2,3,...\infty \\ n$$

- R = alkyl, such as C_nH_{2n+1}, wherein n = 1 to 20, aryl, such as phenyl und substituted phenyl radicals, (CH₂)_n-NH₂, H
- R' = alkyl, such as C_nH_{2n+1} , wherein n = 1 to 20, aryl, such as phenyl- and substituted phenyl radicals, $(CH_2)_n$ -NH2, H
- R' = alkyl, such as C_nH_{2n+1}, wherein n = 1 to 20, aryl, such as phenyl- and substituted phenyl radicals, (CH₂)_n-NH₂, H
- R' = alkyl, such as C_nH_{2n+1}, wherein n = 1 to 20, aryl, such as phenyl und substituted phenyl radicals, (CH₂)_n·NH₂, H₂

2. (Cancelled)

3. (Previously presented) A process for the preparation of the surface-modified pyrogenically produced zinc oxide powder according to Claim 1, comprising optionally spraying a zinc oxide with water, spraying the surface-modifying agent at room temperature to obtain a zinc oxide sprayed with said surface-modifying agent, heat treating said zinc oxide at a temperature of 50 to 400°C over a period of 1 to 6 hours to thereby obtain the surface-modified zinc oxide.

4. (Cancelled)

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5. (Previously presented) A process for the preparation of the surface-modified pyrogenically produced zinc oxide powder according to Claim 1, comprising optionally spraying zinc oxide with water, treating said zinc oxide with the surface-modifying agent in vapour form and then heat-treating the resulting zinc oxide at a temperature of 50 to 800°C over a period of 0.5 to 6 hours to thereby obtain the surface-modified zinc oxide.

6. (Cancelled)

7. (Previously presented) A cosmetic preparation comprising a dermatologically acceptable carrier and the surface-modified pyrogenically produced zinc oxide powder of Claim 1.

8. (Cancelled)

9. (Previously presented) A sunscreen preparation comprising a dermatologically acceptable carrier and the surface modified pyrogenically produced zinc oxide powder of Claim 1.

10. (Cancelled)

11. (Previously presented) The sunscreen preparation according to Claim 9, wherein the dermatologically acceptable carrier is a member selected from the group consisting of octocrylene, ethylhexyl methoxycinnamate, phenylbenzimidazole sulfonic acid, and bisethylhexyloxy methoxyphenyl triazine.

12. (Cancelled)

13. (Cancelled)

14. (Previously presented) The surface-modified pyrogenically produced zinc oxide powder according to Claim 1 wherein the zinc oxide powder displays at its surface an oxygen concentration as non-desorbable moisture in the form of Zn-OH and/or Zn-OH₂ units of at least 40%.